# Title: Is there a relationship between chocolate consumption and symptoms of depression? A cross-sectional survey of 13,626 US adults

**Running title**: Chocolate consumption and depressive symptoms

Sarah E Jackson1, Lee Smith2, Joseph Firth3, Igor Grabovac4, Pinar Soysal5, Ai Koyanagi6, Liang Hu7, Brendon Stubbs8, Jacopo Demurtas9, Nicola Veronese10, Xiangzhu Zhu\*11, Lin Yang\*12

1Department of Behavioural Science and Health, University College London, London, UK

2Cambridge Center for Exercise Science, Anglia Ruskin University, Cambridge, UK

3 NICM Health Research Institute, Western Sydney University, Westmead, Australia; Division of Psychology and Mental Health, University of Manchester, Manchester, UK; Centre for Youth Mental Health, University of Melbourne, Melbourne, Australia

4 Department of Social and Preventive Medicine, Center for Public Health, Medical University of Vienna, Vienna, Austria

5 Department of Geriatric Medicine, Bezmialem Vakif University, Faculty of Medicine, Istanbul, Turkey

6 Research and Development Unit, Parc Sanitari Sant Joan de Déu, CIBERSAM, Barcelona, Spain; ICREA, Pg. Lluis Companys 23, Barcelona, Spain

7 Department of Sport and Exercise Science, Zhejiang University, Hangzhou, China

8 Physiotherapy Department, South London and Maudsley NHS Foundation Trust, Denmark Hill, London, UK; Department of Psychological Medicine, Institute of Psychiatry, Psychology and Neuroscience, King's College London, De Crespigny Park, London, United Kingdom.

9 Primary Care Department Azienda USL Toscana Sue Est, 58100 Grosseto, Italy.

10 National Research Council, Neuroscience Institute, Aging Branch, Padova, Italy.

11 Division of Epidemiology, Department of Medicine, Vanderbilt University Medical Center, Nashville, USA; Vanderbilt Ingram Cancer Center, Vanderbilt University School of Medicine, Nashville, USA.

12 Cancer Epidemiology and Prevention Research, Alberta Health Services, Calgary, Alberta, Canada; Departments of Oncology and Community Health Sciences, Cumming School of Medicine, University of Calgary, Calgary, Alberta, Canada

\* These authors contribute equally to this work.

**Corresponding authors**: Sarah E Jackson, Ph.D, s.e.jackson@ucl.ac.uk; Lin Yang, Ph.D, lin.yang@ahs.ca

**Acknowledgement**: The authors received no financial support for the research, authorship, and publication of this article.

**Conflict of Interest Statement**: The authors have no conflict of interest to declare.

## Abstract

**Objective:** To examine associations between chocolate consumption and depressive symptoms in a large, representative sample of US adults.

**Methods:** Data were used from 13,626 adults (≥20y) participating in the National Health and Nutrition Examination Survey between 2007-08 and 2013-14. Daily chocolate consumption was derived from two 24-hour dietary recalls. Depressive symptoms were assessed using the Patient Health Questionnaire (PHQ-9), with scores ≥10 indicating the presence of clinically relevant symptoms. We used multivariable logistic regression to test associations of chocolate consumption (no chocolate, non-dark chocolate, dark chocolate) and amount of chocolate consumption (grams/day, in quartiles) with clinically relevant depressive symptoms. Adults with diabetes were excluded and models controlled for relevant sociodemographic, lifestyle, health-related, and dietary covariates.

**Results:** Overall, 11.1% of the population reported any chocolate consumption, with 1.4% reporting dark chocolate consumption. Although non-dark chocolate consumption was not significantly associated with clinically relevant depressive symptoms, significantly lower odds of clinically relevant depressive symptoms (OR=0.30, 95%CI 0.21-0.72) were observed among those who reported consuming dark chocolate. Analyses stratified by amount of chocolate consumption showed participants reporting chocolate consumption in the highest quartile (104-454g/day) had 57% lower odds depressive symptoms than those who reported no chocolate consumption (OR=0.43, 95%CI 0.19-0.96) after adjusting for dark chocolate consumption.

**Conclusions:** These results provide some evidence that consumption of chocolate, particularly dark chocolate, may be associated with reduced odds of clinically relevant depressive symptoms. Further research capturing long-term chocolate consumption and using a longitudinal design are required to confirm these findings and clarify the direction of causation.

**Keywords**: Chocolate, dark chocolate, depressive symptoms, Epidemiology, NHANES

## Introduction

Depression is a serious, common, and recurring disorder that currently affects more than 300 million people worldwide (World Health Organization, 2018). It is the leading cause of disability and the fourth leading cause of global disease burden (Ustun, Ayuso-Mateos, Chatterji, Mathers, & Murray, 2004). Two main treatment options for depression have been shown to be effective in reducing symptoms: antidepressant medications and short-term psychotherapies, such as cognitive behavioural therapy, interpersonal psychotherapy, or problem-solving therapy (Mynors-Wallis, Gath, Day, & Baker, 2000; Schulberg et al., 1996; Ward et al., 2000). Due to limited availability of psychotherapy services, antidepressants are generally the most commonly prescribed treatment for depression in primary care (van marwijk, Bijl, Ader, & de Haan, 2001). However, patient adherence is relatively low, with as many as half of patients prescribed these medications discontinuing their use within six weeks of starting treatment (Lawrenson, Tyrer, Newson, & Farmer, 2000).

The potential to control depressive symptoms via lifestyle changes may provide an attractive option to patients for whom conventional treatments are not an effective option and individuals with subclinical depressive symptoms who are not eligible for formal treatment. Benefits of physical activity are well documented (Schuch et al., 2018; Stubbs et al., 2018; Teychenne, Ball, & Salmon, 2008). In addition, a growing body of evidence has highlighted the influence of dietary factors on symptoms of depression (Molendijk, Molero, Ortuno Sanchez-Pedreno, Van der Does, & Angel Martinez-Gonzalez, 2018). One commonly consumed foodstuff postulated to have mood-enhancing properties is chocolate. Several mechanisms for a relationship between chocolate and mood have been proposed, including chocolate’s orosensory properties, psychoactive ingredients, and activation of neural reward pathways (Bruinsma & Taren, 1999; Hurst, Martin, & Zoumas, 1982; Parker, Parker, & Brotchie, 2006). However, there is a lack of high-quality scientific evidence to support such an association. Only a small number of studies have analysed associations between chocolate consumption and depressive symptoms, with mixed findings. A 2013 systematic review of experimental research on the effects of chocolate or its components on mood identified eight small studies (*n*≤113), of which five reported either an improvement in mood state or an attenuation of negative mood (Scholey & Owen, 2013). A recent umbrella review on chocolate consumption and all health outcomes concluded based on existing evidence that chocolate consumption has no impact on depression (Veronese et al., 2018). A cross-sectional survey of 931 adults without diabetes or known coronary artery disease found higher depressive symptom scores were associated with greater chocolate consumption in both men and women (Rose, Koperski, & Golomb, 2010). Likewise, another study of 362 women observed a positive correlation between reported chocolate consumption and scores on a measure of psychiatric symptomology (Barkeling et al., 2002). However, previous studies have not adequately controlled for variables that may potentially confound the association between chocolate and depression, such as socioeconomic status.

Moreover, previous studies have not examined the association with depression according to the type of chocolate consumed. There is a large body of literature demonstrating a positive association between dark chocolate consumption and physical health outcomes (Mursu et al., 2004; Tokede, Gaziano, & Djousse, 2011; Vlachopoulos et al., 2005). This beneficial effect of dark chocolate is likely owing to flavonoids that have antioxidant properties and improve inflammatory profiles (Minihane et al., 2015). Importantly, flavonoids have also been shown to have a positive influence on episodic memory and cognitive decline (Minihane et al., 2015). It may therefore be that dark chocolate (rich in flavonoids) has a positive effect on mental health, including depression, whereas milk chocolate (lower in flavonoids) does not. However, to date the association between consumption of dark chocolate and depression has not been investigated.

The present study therefore aimed to examine the relationship between chocolate consumption and symptoms of depression in a large, nationally-representative sample of adults living in the US, with adjustment for a wide range of potential confounders. Specifically, we were interested in: (i) the association between chocolate consumption (any dark chocolate, chocolate but not dark chocolate, none) and depressive symptoms among all adults, (ii) the association between amount of chocolate consumption and depressive symptoms.

## Materials and methods

### Study population

The National Health and Nutrition Examination Survey (NHANES), described in detail elsewhere (Curtin et al., 2013, 2012), was designed to provide cross-sectional estimates of the prevalence of health, nutrition, and potential risk factors among the civilian non-institutionalized US population (Centers for Disease Control and Prevention). Since 1999, NHANES has surveyed a nationally representative, complex, stratified, multistage probability sample of the US population continuously in 2-year cycles, with different participants included in each wave. The assessment methods include a household interview and a physical examination in a mobile examination centre (MEC). For the present analyses, we included participants over five cycles from 2007-08 to 2013-14 with available data on chocolate consumption from two 24-hour dietary recalls. We restricted our study sample to adults aged ≥20 years because the assessment of depressive symptoms was only done in this age range.

### Measures

#### Exposure: chocolate consumption

A key component of NHANES is dietary assessment. In brief, NHANES participants are asked to complete two 24-hour recalls of dietary intake using the USDA’s Automated Multiple-Pass method (Ahluwalia, Dwyer, Terry, Moshfegh, & Johnson, 2016). For the cycles used in the present analyses, the first 24-hour recall was administered in person by a trained dietary interviewer using a standardized protocol during the physical examination in a mobile examination center. The second 24-hour recall was administered via telephone between 3 and 10 days after the first recall. Nutrient intakes were calculated based on food intake using a revised nutrient database that converts intakes of food items to nutrients for each individual. Daily chocolate consumption (overall and dark chocolate [≥45% cocoa solids] specifically) was calculated in grams using the average of the two 24-hour recalls, with implausibly high intakes excluded by removing observations above the 99th centile. The present analyses focus on two variables derived from this measure of chocolate consumption. The first was a three-level categorical measure of chocolate consumption (none, chocolate but not dark chocolate, any dark chocolate). The second was a five-level measure of the amount of chocolate consumed (any type) in gram/day. Individuals who reported no chocolate consumption were coded 0, and those who reported any chocolate consumption were grouped into quartiles, coded 1 through 4 from the lowest to highest chocolate consumption.

#### Outcome: depressive symptoms

Depressive symptoms were assessed using the Patient Health Questionnaire (PHQ-9), a valid 9-item depression screener that asks about the frequency of symptoms of depression over the past two weeks (Kroenke, Spitzer, & Williams, 2001; Mitchell, Yadegarfar, Gill, & Stubbs, 2016). Each item was scored on a 0-3 scale, with a total score ranging from 0 to 27. On the basis of these scores, depressive symptoms can be categorised as “none or minimum” (0-4), “mild” (5-9), “moderate” (10-14), “moderately severe” (15-19), and “severe” (20-27). For the present analyses, the data were dichotomized to distinguish between participants who scored 9 or less and those who scored ≥10 or more; the latter indicating clinically relevant depression. This cut-off has previously been shown to have a sensitivity of 88% and a specificity of 88% for major depression (Kroenke et al., 2001; Manea, Gilbody, & McMillan, 2012; Mitchell et al., 2016).

#### Covariates

Total energy intake and total sugar intake were derived from the mean of the two 24-hour dietary recalls, and categorized into quartiles. Alcohol intake was derived from two 24-hour recalls and dichotomized to no (zero intake) and yes (any intake).

Weight and height were measured during the physical examination in a mobile examination center or in the participant’s home, following standard procedures. Body mass index (BMI) was calculated as weight in kilograms divided by the square of height in meters and categorized as underweight (<18.5 kg/m2), normal weight (18.5 – 24.9 kg/m2), overweight (25.0 – 29.9 kg/m2), or obese (≥30 kg/m2). For analytic purposes, we excluded those who with an underweight BMI due to potential underlying health conditions.

Self-reported socio-demographic characteristics included age, marital status (collapsed to living alone [widowed, divorced, separated, never married] and living with someone [married, living with partner]), sex (men and women), race (non-Hispanic white, non-Hispanic black, Hispanic, other [other than non-Hispanic white, non-Hispanic black, Hispanic, including multi-racial]), education (collapsed to below high school [less than 9th grade, 9-11th grade], high school [high school graduate/GED or equivalent], above high school [some college or AA degree, college graduate or above]) and annual household income (15 categories of annual household income levels collapsed to <$25,000, $25,000–$74,999, and ≥$75,000).

Lifestyle characteristics included leisure-time physical activity and smoking status. Participants reported the number of days and minutes spent in moderate and vigorous recreational activities in a typical week. We summarized the total number of minutes for both activities and classified participants as inactive (zero moderate-to-vigorous physical activity) or active (any moderate-to-vigorous physical activity) based on physical activity guidelines (U.S. Department of Health and Human Services, 2018). Smoking status was classified into: never smokers (have never smoked 100 cigarettes and do not smoke now), former smokers (have smoked 100 cigarettes in lifetime and do not smoke now), and current smokers (have smoked 100 cigarettes in lifetime and smoke now).

Information on chronic conditions that were considered suspected correlates were extracted, including self-reported doctors’ diagnoses (by answering the question “Has a doctor or other health professional ever told you that you had…” for each condition) of cardiovascular disease, arthritis, and cancer. Participants who reported diabetes mellitus were excluded from the analyses due to potential confounding by increased prevalence of depression (Roy & Lloyd, 2012) and likelihood of dietary restrictions that prohibit chocolate consumption.

### Statistical analyses

All statistical analyses were performed using STATA version 14.0 (STATA Corp., College Station, Texas, USA). Survey analysis procedures were used to account for the sample weights, stratification, and clustering of the complex sampling design to ensure nationally representative estimates (Curtin et al., 2012). Descriptive characteristics were analysed by summarizing weighted means and standard errors for continuous variables and weighted proportions for categorical variables by chocolate consumption status (none, non-dark chocolate, dark chocolate) as well as by quartiles of chocolate consumption.

Three multivariable logistic regression models were constructed to evaluate the association of chocolate consumption with clinically relevant depressive symptoms. First, we estimated the independent association of chocolate consumption (none, non-dark chocolate, and dark chocolate) with clinically relevant depressive symptoms in the entire sample. Secondly, we tested whether the amount of chocolate consumption (zero for non-chocolate consumers, and in quartiles among chocolate consumers) was independently associated with clinically relevant depressive symptoms, with and without adjustment for dark chocolate consumption (yes/no). It was not possible to analyse the association with the amount of dark chocolate consumption specifically due to the small number of participants reporting any dark chocolate consumption. All models were adjusted for sociodemographic factors (age, sex, marital status, education level, annual household income), weight status, lifestyle factors (leisure-time physical activity, smoking status, alcohol intake, total energy intake, and chronic conditions. We constructed models with and without total sugar intake to assess whether the adjustment for sugar consumption may lead to over-adjustment, as there might be a potential overlap between chocolate consumption and sugar intake. Diagnostic tests were performed to evaluate the degree of multicollinearity in each model. We considered *p* values <0.05 to be statistically significant.

## Results

A total of 20,125 men and women aged ≥20 years provided data on chocolate consumption and depressive symptoms. We excluded those who were underweight (n=304), and those who with diabetes (n=2518). A further 3,677 participants were excluded due to missing covariates, leaving a total of 13,626 in the final analyses (mean age 46.5 years, 47.8% male). The individuals we excluded had similar prevalence of clinically relevant depressive symptoms to the analysed sample (Chi-square p=0.18), but there were more chocolate consumers in the analysed sample (11.1% vs. 7.0%, Chi-square p<.001).

Of our analysed sample, 1,332 (11.1%) adults reported any chocolate consumption in their two 24-hour dietary recalls, of whom 148 (12.1%) reported any dark chocolate consumption. Sample characteristics in relation to chocolate consumption are summarised in Table 1 (for sample characteristics in relation to quartiles of chocolate consumption see supplemental table 1). Those who reported any chocolate consumption were slightly older on average than those who did not report chocolate consumption. They were more likely to be non-Hispanic white and have a higher household income and level of education, and were less likely to smoke or have a BMI in the obese range. Average daily intakes of total energy and sugar were higher among adults who were non-dark chocolate consumers. In addition, chocolate consumption was not significantly associated with alcohol drinking nor chronic conditions in our study population.

After adjustment for covariates, no association was observed between non-dark chocolate consumption and clinically relevant depressive symptoms (Table 2). However, individuals who reported any dark chocolate consumption had 70% lower odds of reporting clinically relevant depressive symptoms than those who did not report any chocolate consumption (OR=0.30, 95% CI 0.21 to 0.72). In models including the amount of chocolate consumption, significant association between the highest quartile of total chocolate consumption and clinically relevant depressive symptoms (OR=0.42, 95% CI: 0.19 to 0.92). This association remained significant after adjusting for dark chocolate consumption (yes/no) (OR=0.43, 95% CI: 0.19 to 0.96) (Table 2). The results were similar with and without the adjustment for sugar intake. Hence, the adjustment for sugar intake is unlikely to lead to an over-adjustment.(Table 2). Furthermore, we have carried out logistic regression diagnostic test for multicollinearity. No severe multicollinearity was observed, with variance inflation factors between 1.03 and 1.93, and tolerance between 0.5172 and 0.9754.

## Discussion

In a large, representative sample of US adults, individuals who reported any consumption of dark chocolate were less likely to report clinically relevant depressive symptoms, but no such association was observed for non-dark chocolate. However, we also observed a significant association with the amount of chocolate consumed, with individuals in the upper quintile of chocolate consumption (calculated among chocolate consumers) significantly less likely to report clinically relevant depressive symptoms than those who reported no chocolate consumption, even after adjustment for type of chocolate. These associations were evident after adjustment for age, marital status, level of education, annual household income, weight status, chronic conditions, leisure-time physical activity, smoking status, alcohol intake, total energy intake, and total sugar intake.

The present results are in line with the majority of experimental studies, which have shown benefits of chocolate consumption for mood, at least in the short-term (Scholey & Owen, 2013). However, they are inconsistent with previous surveys that have found positive associations between chocolate consumption and depressive symptoms (Barkeling et al., 2002; Rose et al., 2010). The discrepant results may be attributable to the adjustment in the present analyses for a wide range of covariates accounting for potential confounding. Alternatively, the representative sample here may differ from those of previous studies, which focused on adults without specific medical conditions (Rose et al., 2010) or women aged 34-64 years (Barkeling et al., 2002).

There are a number of potential mechanisms through which chocolate consumption could prevent the onset of, or cause a reduction in, depressive symptoms. Chocolate contains a number of psychoactive ingredients, including two analogues of anandamine (which produce effects similar to the cannabinoid responsible for euphoria from cannabis) and several endogenous biogenic amines (Bruinsma & Taren, 1999; Hurst et al., 1982; Parker et al., 2006). Notably, the latter include phenylethylamine, a neuromodulator believed to be important for mood regulation and implicated in the pathogenesis of depression (Bruinsma & Taren, 1999; Sabelli & Javaid, 1995). Chocolate consumption, as a pleasurable experience, may interact with several neurotransmitter systems implicated in reward pathways and mood regulation (e.g. dopamine, serotonin, and endorphins) (Bruinsma & Taren, 1999; Parker et al., 2006). Experimental evidence suggests that improvements in mood following chocolate consumption is likely attributable to the chocolate’s palatability (Macht & Dettmer, 2006; Macht & Mueller, 2007). In a study that compared effects of eating palatable chocolate, unpalatable chocolate, and nothing on experimentally-induced negative mood states, an immediate benefit was only seen after consumption of palatable chocolate, with no difference between unpalatable chocolate and nothing (Macht & Mueller, 2007). However, this would not explain why the associations observed in the present study were more pronounced for dark chocolate than for all chocolate. It is possible that the rich concentration of flavonoids in dark chocolate may be important. Indeed, flavonoids have been shown to improve in inflammatory profiles and unfavourable inflammatory profiles have been shown to play a role in the onset of depression (Penninx, Milaneschi, Lamers, & Vogelzangs, 2013). Further research is required to elucidate the nature of the relationship between chocolate consumption and depressive symptoms.

Strengths of the present study include the large, nationally-representative sample, detailed dietary assessment, and adjustment for a wide range of relevant covariates. However, there were also several limitations. First, the cross-sectional design means we were unable to determine whether chocolate consumption protected against the onset of depressive symptoms. An alternative explanation may be depressive symptoms make people less inclined to eat chocolate. However, previous research has suggested that almost half (44.9%) of people who have experienced depressive episodes lasting two weeks or longer that required treatment report craving chocolate when depressed, with many believing that chocolate helps to settle feelings of anxiety and irritability (Parker & Crawford, 2007). Secondly, chocolate consumption was based on the average of two 24-hour dietary recalls. While dietary recalls are generally considered to provide a more robust measure of nutrient consumption than food frequency questionnaires, which ask participants to report how often they eat different types of foods, the measure of chocolate consumption in the present study may not accurately reflect average intake over a prolonged period because chocolate is not a product that is typically consumed every day. As such, the potential for misclassification of chocolate consumption is high. There is a need to replicate our findings using other measures of chocolate consumption that capture usual intake over the longer term. Thirdly, the group of dark chocolate consumers in our analyzed sample was small with low consumption (less than 1.1% of the total population with an average of 11.7 grams of daily dark chocolate intake) and differed from the rest of the population on a number of key characteristics. Nonetheless, the fact we observed a significant association between dark chocolate consumption despite the small sample and low average consumption attests to the strength of this finding. Although we did adjust for a number of covariates, it is possible that we lacked information on some characteristics of dark chocolate consumers that could have confounded the associations assessed. Thus, some caution should be taken when interpreting these results. Finally, while we adjusted for a wide range of potential confounders, there may be residual confounding by unmeasured variables. Studies using a randomized design to test the effect of chocolate consumption (in particular, dark chocolate) on mood over a prolonged period could provide further insight into the exact nature of the relationship between these variables.

In conclusion, the present results provide some evidence that consumption of chocolate, particularly dark chocolate, may be associated with reduced odds of clinically relevant depressive symptoms. Further research is required to clarify the direction of causation. Should a causal relationship demonstrating a protective effect of chocolate consumption on depressive symptoms be established, its biological mechanism needs to be elucidated to determine the type and amount of chocolate consumption for optimal depression prevention and management.

**Data Availability Statement**:The NHANES data that support the findings of this study are available online through https://wwwn.cdc.gov/nchs/nhanes/Default.aspx

## References

Ahluwalia, N., Dwyer, J., Terry, A., Moshfegh, A., & Johnson, C. (2016). Update on NHANES Dietary Data: Focus on Collection, Release, Analytical Considerations, and Uses to Inform Public Policy. *Adv Nutr, 7*(1), 121-134. doi:10.3945/an.115.009258

Barkeling, B., Linne, Y., Lindroos, A. K., Birkhed, D., Rooth, P., & Rossner, S. (2002). Intake of sweet foods and counts of cariogenic microorganisms in relation to body mass index and psychometric variables in women. *Int J Obes Relat Metab Disord, 26*(9), 1239-1244. doi:10.1038/sj.ijo.0802034

Bruinsma, K., & Taren, D. L. (1999). Chocolate: food or drug? *J Am Diet Assoc, 99*(10), 1249-1256. doi:10.1016/s0002-8223(99)00307-7

Centers for Disease Control and Prevention. NHANES - National Health and Nutrition Examination Survey Homepage [Internet]. 2018. Available from: https://www.cdc.gov/nchs/nhanes/index.htm accessed March 19, 2018.

Curtin, L. R., Mohadjer, L. K., Dohrmann, S. M., Montaquila, J. M., Kruszan-Moran, D., Mirel, L. B., . . . Johnson, C. L. (2012). The National Health and Nutrition Examination Survey: Sample Design, 1999-2006. *Vital Health Stat 2*(155), 1-39.

Curtin, L. R., Mohadjer, L. K., Dohrmann, S. M., Kruszon-Moran, D., Mirel, L. B., Carroll, M. D., … Johnson, C. L. (2013). National Health and Nutrition Examination Survey: sample design, 2007-2010. *Vital and Health Statistics. Series 2, Data Evaluation and Methods Research*, (160), 1–23.

Hurst, W. J., Martin, R. A., & Zoumas, B. L. (1982). Biogenic amines in chocolate--a review. *Nutr Rep Int, 26*, 1081-1086.

Kroenke, K., Spitzer, R. L., & Williams, J. B. (2001). The PHQ-9: validity of a brief depression severity measure. *J Gen Intern Med, 16*(9), 606-613.

Lawrenson, R. A., Tyrer, F., Newson, R. B., & Farmer, R. D. (2000). The treatment of depression in UK general practice: selective serotonin reuptake inhibitors and tricyclic antidepressants compared. *J Affect Disord, 59*(2), 149-157.

Macht, M., & Dettmer, D. (2006). Everyday mood and emotions after eating a chocolate bar or an apple. *Appetite, 46*(3), 332-336. doi:10.1016/j.appet.2006.01.014

Macht, M., & Mueller, J. (2007). Immediate effects of chocolate on experimentally induced mood states. *Appetite, 49*(3), 667-674. doi:10.1016/j.appet.2007.05.004

Manea, L., Gilbody, S., & McMillan, D. (2012). Optimal cut-off score for diagnosing depression with the Patient Health Questionnaire (PHQ-9): a meta-analysis. *Cmaj, 184*(3), E191-196. doi:10.1503/cmaj.110829

Minihane, A. M., Vinoy, S., Russell, W. R., Baka, A., Roche, H. M., Tuohy, K. M., . . . Calder, P. C. (2015). Low-grade inflammation, diet composition and health: current research evidence and its translation. *Br J Nutr, 114*(7), 999-1012. doi:10.1017/s0007114515002093

Mitchell, A. J., Yadegarfar, M., Gill, J., & Stubbs, B. (2016). Case finding and screening clinical utility of the Patient Health Questionnaire (PHQ-9 and PHQ-2) for depression in primary care: a diagnostic meta-analysis of 40 studies. *BJPsych Open, 2*(2), 127-138. doi:10.1192/bjpo.bp.115.001685

Molendijk, M., Molero, P., Ortuno Sanchez-Pedreno, F., Van der Does, W., & Angel Martinez-Gonzalez, M. (2018). Diet quality and depression risk: A systematic review and dose-response meta-analysis of prospective studies. *J Affect Disord, 226*, 346-354. doi:10.1016/j.jad.2017.09.022

Mursu, J., Voutilainen, S., Nurmi, T., Rissanen, T. H., Virtanen, J. K., Kaikkonen, J., . . . Salonen, J. T. (2004). Dark chocolate consumption increases HDL cholesterol concentration and chocolate fatty acids may inhibit lipid peroxidation in healthy humans. *Free Radic Biol Med, 37*(9), 1351-1359. doi:10.1016/j.freeradbiomed.2004.06.002

Mynors-Wallis, L. M., Gath, D. H., Day, A., & Baker, F. (2000). Randomised controlled trial of problem solving treatment, antidepressant medication, and combined treatment for major depression in primary care. *Bmj, 320*(7226), 26-30.

Parker, G., & Crawford, J. (2007). Chocolate craving when depressed: a personality marker. *Br J Psychiatry, 191*, 351-352. doi:10.1192/bjp.bp.106.033746

Parker, G., Parker, I., & Brotchie, H. (2006). Mood state effects of chocolate. *J Affect Disord, 92*(2-3), 149-159. doi:10.1016/j.jad.2006.02.007

Penninx, B. W., Milaneschi, Y., Lamers, F., & Vogelzangs, N. (2013). Understanding the somatic consequences of depression: biological mechanisms and the role of depression symptom profile. *BMC Med, 11*, 129. doi:10.1186/1741-7015-11-129

Rose, N., Koperski, S., & Golomb, B. A. (2010). Mood food: chocolate and depressive symptoms in a cross-sectional analysis. *Arch Intern Med, 170*(8), 699-703. doi:10.1001/archinternmed.2010.78

Roy, T., & Lloyd, C. E. (2012). Epidemiology of depression and diabetes: a systematic review. *J Affect Disord, 142 Suppl*, S8-21. doi:10.1016/s0165-0327(12)70004-6

Sabelli, H. C., & Javaid, J. I. (1995). Phenylethylamine modulation of affect: therapeutic and diagnostic implications. *J Neuropsychiatry Clin Neurosci, 7*(1), 6-14. doi:10.1176/jnp.7.1.6

Scholey, A., & Owen, L. (2013). Effects of chocolate on cognitive function and mood: a systematic review. *Nutr Rev, 71*(10), 665-681. doi:10.1111/nure.12065

Schuch, F. B., Vancampfort, D., Firth, J., Rosenbaum, S., Ward, P. B., Silva, E. S., … Stubbs, B. (2018). Physical Activity and Incident Depression: A Meta-Analysis of Prospective Cohort Studies. Th*e American Journal of Psychiatry*, 175(7), 631–648. https://doi.org/10.1176/appi.ajp.2018.17111194

Schulberg, H. C., Block, M. R., Madonia, M. J., Scott, C. P., Rodriguez, E., Imber, S. D., . . . Coulehan, J. L. (1996). Treating major depression in primary care practice. Eight-month clinical outcomes. *Arch Gen Psychiatry, 53*(10), 913-919.

Stubbs, B., Vancampfort, D., Hallgren, M., Firth, J., Veronese, N., Solmi, M., … Kahl, K. G. (2018). EPA guidance on physical activity as a treatment for severe mental illness: a meta-review of the evidence and Position Statement from the European Psychiatric Association (EPA), supported by the International Organization of Physical Therapists in Mental Health (IOPTMH). European Psychiatry: *The Journal of the Association of European Psychiatrists*, 54, 124–144. https://doi.org/10.1016/j.eurpsy.2018.07.004

Teychenne, M., Ball, K., & Salmon, J. (2008). Physical activity and likelihood of depression in adults: a review. *Prev Med, 46*(5), 397-411. doi:10.1016/j.ypmed.2008.01.009

Tokede, O. A., Gaziano, J. M., & Djousse, L. (2011). Effects of cocoa products/dark chocolate on serum lipids: a meta-analysis. *Eur J Clin Nutr, 65*(8), 879-886. doi:10.1038/ejcn.2011.64

U.S. Department of Health and Human Services. (2018). *Physical Activity Guidelines for Americans, 2nd edition. Washington DC: U.S. Department of Health and Human Services*. Retrieved from

Ustun, T. B., Ayuso-Mateos, J. L., Chatterji, S., Mathers, C., & Murray, C. J. (2004). Global burden of depressive disorders in the year 2000. *Br J Psychiatry, 184*, 386-392.

van Marwijk, H. W., Bijl, D., Ader, H. J., & de Haan, M. (2001). Antidepressant prescription for depression in general practice in The Netherlands. *Pharm World Sci, 23*(2), 46-49.

Veronese, N., Demurtas, J., Celotto, S., Caruso, M. G., Maggi, S., Bolzetta, F., . . . Stubbs, B. (2018). Is chocolate consumption associated with health outcomes? An umbrella review of systematic reviews and meta-analyses. *Clin Nutr*. doi:10.1016/j.clnu.2018.05.019

Vlachopoulos, C., Aznaouridis, K., Alexopoulos, N., Economou, E., Andreadou, I., & Stefanadis, C. (2005). Effect of dark chocolate on arterial function in healthy individuals. *Am J Hypertens, 18*(6), 785-791. doi:10.1016/j.amjhyper.2004.12.008

Ward, E., King, M., Lloyd, M., Bower, P., Sibbald, B., Farrelly, S., . . . Addington-Hall, J. (2000). Randomised controlled trial of non-directive counselling, cognitive-behaviour therapy, and usual general practitioner care for patients with depression. I: clinical effectiveness. *Bmj, 321*(7273), 1383-1388.

World Health Organization. Depression [Internet]. 2018. Available from: https://www.who.int/news-room/fact-sheets/detail/depressionDemographisches accessed March 19, 2018.

|  |
| --- |
| Table 1. Socio-demographic characteristics of US adults aged 20-80 Years in the NHANES (2007-2014), by chocolate consumption statusa |
|   |   | Chocolate consumption |
|  |   | None | Non-dark chocolate | Dark chocolate | P-value |
|  |   | (n=12,294) | (n=1,184) | (n=148) |   |
| Age (years) | mean (se) | 46.2(0.3) | 47.8 (0.5) | 57.3 (1.2) | <0.001 |
| Gender (men) | % | 48.8 | 39.7 | 43.8 |  |
| Marital status (living with someone) | % | 64.3 | 68.1 | 74.0 | <0.001 |
| Race |  |  |  |  | <0.001 |
| Non-Hispanic White | % | 70.5 | 80.2 | 86.7 |  |
| Non-Hispanic Black | % | 10.6 | 4.4 | 3.0 |  |
| Hispanic  | % | 13.0 | 9.6 | 4.8 |  |
| Other | % | 5.9 | 5.8 | 5.5 |  |
| Household income |  |  |  | <0.001 |
| ≤ $20000 | % | 20.6 | 15.7 | 11.0 |  |
| $20000-74999 | % | 42.1 | 38.7 | 34.4 |  |
| ≥$75000 | % | 37.3 | 45.6 | 54.7 |  |
| Education |  |  |  | <0.001 |
| Less than High school | % | 15.4 | 10.1 | 4.1 |  |
| High School | % | 22.3 | 19.0 | 15.3 |  |
| Some college and above | % | 62.4 | 70.9 | 80.6 |  |
| Smoking |  |  |  |  | <0.001 |
| Never smoker | % | 55.6 | 61.5 | 64.1 |  |
| Former smoker | % | 24.1 | 24.7 | 26.9 |  |
| Current smoker | % | 20.3 | 13.8 | 9.0 |  |
| Alcohol drinking (yes) | % | 36.1 | 36.9 | 46.3 | 0.39 |
| BMI (kg/m2) |  |  |  | <0.001 |
| 18.5 – 24.9 | % | 29.6 | 34.9 | 38.8 |  |
| 25.0 – 29.9  | % | 35.2 | 39.2 | 32.9 |  |
| ≥ 30 | % | 35.2 | 25.9 | 28.3 |  |
| Chronic conditions (yes)b | % | 28.8 | 32.3 | 42.0 | 0.004 |
| Depressive symptoms (yes)c | % | 7.6 | 6.2 | 1.5 | 0.019 |
| Total energy intake (kcal, day) | mean (se) | 2115.6 (10.5) | 2295.7 (35.7) | 2122.6 (88.2) | 0.014 |
| Total sugar intake (gram, day) | mean (se) | 113.2 (0.9) | 127.6 (3.0) | 120.2 (6.4) | <0.001 |
| Total chocolate intake (gram, day) | mean (se) | n.a | 91.8 (2.7) | 77.7 (5.8) |  |
| Total dark chocolate intake (gram, day) | mean (se) | n.a | n.a | 11.7 (1.0) |  |
| a. All estimated are weighted to be nationally representative |
| b. Depression symptoms measured using Patient Health Questionnaire (PHQ-9): None or minimum to mild (0-9) Moderate to severe (10-27) |
| c. Chronic conditions include cardiovascular disease, arthritis, and cancer. |

|  |
| --- |
| Table 2 Associations of chocolate consumption with clinically relevant depressive symptoms among adults aged 20-80 years (n=13,626) in NHANES (2007-2014)a |
|  | Unadjusted |  | Multivariable-adjustedb |  | Multivariable-adjustedc |  | Multivariable-adjustedd |
|  | OR(95% CI) |  | OR(95% CI) |  | OR(95% CI) |  | OR(95% CI) |
| Overall chocolateb |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| None | Reference |  |  | Reference |  |  | Reference |  |  |  |  |
| Non-dark chocolate | 0.80 | (0.59 to | 1.08) |  | 0.95 | (0.70 to | 1.28) |  | 0.94 | (0.69 to | 1.27) |  |  |  |  |
| Dark chocolate | 0.19 | (0.07 to | 0.48) |  | 0.30 | (0.12 to | 0.72) |  | 0.30 | (0.12 to | 0.72) |  |  |  |  |
| Daily chocolate consumptionb |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| None | Reference |  |  | Reference |  |  | Reference |  |  | Reference |  |
| 1st quartile (4.1 – 35.0 grams) | 0.95 | (0.53 to | 1.71) |  | 1.03 | (0.57 to | 1.87) |  | 1.03 | (0.57 to | 1.86) |  | 1.03 | (0.57 to | 1.87) |
| 2nd quartile (37.0 – 95.1 grams) | 0.45 | (0.24 to | 0.84) |  | 0.56 | (0.30 to | 1.03) |  | 0.55 | (0.30 to | 1.02) |  | 0.64 | (0.33 to | 1.22) |
| 3rd quartile (100 – 100 grams) | 0.98 | (0.61 to | 1.57) |  | 1.28 | (0.80 to | 2.03) |  | 1.27 | (0.79 to | 2.02) |  | 1.29 | (0.80 to | 2.07) |
| 4th quartile (104 – 454 grams) | 0.34 | (0.16 to | 0.73) |  | 0.42 | (0.19 to | 0.93) |  | 0.42 | (0.19 to | 0.92) |  | 0.43 | (0.19 to | 0.96) |
| P for trende | 0.015 |  | 0.224 |  | 0.211 |  | 0.325 |
| a. Depression symptoms measured using Patient Health Questionnaire (PHQ-9): None or minimum to mild (0-9) Moderate to severe (10-27) |
| b. Adjusted for age, sex, BMI category (normal weight, over weight, obese), race (non-Hispanic white, non-Hispanic black, Hispanic, other), education (below high school, high school, high school and higher), household annual income (<$20000, $20000-74999, $75000 and higher), physical activity (inactive, any activity), smoking status (never smoker, former smoker, current smoker), energy intake in quartiles, sugar intake in quartiles, alcohol consumption (yes/no), and chronic conditions including cardiovascular disease, arthritis and cancer (yes/no). |
| c. Additionally adjusted for daily sugar intake (gram, day) |
| d. Additionally adjusted for dark chocolate consumption (yes/no). |
| e. P for trend was calculated by modeling the median value of chocolate consumption in each quartile as a continuous variable. |

|  |
| --- |
| Supplemental Table 1. Socio-demographic characteristics of US adults aged 20-80 Years in the NHANES (2007-2014), by daily chocolate consumptiona |
|   |   | Chocolate consumption |
|  |   | None | 1st quartile(4.1 – 35.0 grams) | 2nd quartile (37.0 – 95.1 grams) | 3rd quartile (100 – 100 grams) | 4th quartile (104 – 454 grams) |
|  |   | (n=12,294) | (n=320) | (n=333) | (n=429) | (n=250) |
| Age (years) | mean (se) | 46.2(0.3) | 47.1 (0.8) | 49.6 (1.2) | 50.7 (0.8) | 47.6 (1.4) |
| Gender (men) | % | 48.8 | 44.2 | 43.4 | 34.9 | 40.7 |
| Marital status (living with someone) | % | 64.3 | 66.1 | 69.7 | 70.1 | 68.5 |
| Race |  |  |  |  |  |  |
| Non-Hispanic White | % | 70.5 | 81.4 | 77.1 | 82.8 | 82.0 |
| Non-Hispanic Black | % | 10.6 | 5.1 | 5.4 | 4.1 | 2.1 |
| Hispanic  | % | 13.0 | 9.8 | 10.5 | 7.7 | 8.5 |
| Other | % | 5.9 | 3.7 | 7.0 | 5.4 | 7.4 |
| Household income |  |  |  |  |  |
| ≤ $20000 | % | 20.6 | 15.1 | 17.0 | 12.3 | 17.7 |
| $20000-74999 | % | 42.1 | 44.1 | 36.4 | 38.2 | 32.9 |
| ≥$75000 | % | 37.3 | 40.8 | 46.6 | 49.5 | 49.4 |
| Education |  |  |  |  |  |
| Less than High school | % | 15.4 | 14.0 | 8.0 | 9.0 | 6.2 |
| High School | % | 22.3 | 21.2 | 17.6 | 18.6 | 16.2 |
| Some college and above | % | 62.4 | 64.8 | 74.4 | 72.4 | 77.6 |
| Smoking |  |  |  |  |  |  |
| Never smoker | % | 55.6 | 56.7 | 57.8 | 63.9 | 69.3 |
| Former smoker | % | 24.1 | 24.9 | 27.5 | 24.4 | 23.1 |
| Current smoker | % | 20.3 | 18.4 | 14.7 | 11.7 | 7.6 |
| Alcohol drinking (yes) | % | 36.1 | 36.9 | 38.8 | 39.8 | 35.5 |
| BMI (kg/m2) |  |  |  |  |  |
| 18.5 – 24.9 | % | 29.6 | 28.7 | 39.0 | 36.3 | 37.8 |
| 25.0 – 29.9  | % | 35.2 | 43.7 | 35.0 | 38.8 | 35.2 |
| ≥ 30 | % | 35.2 | 27.6 | 26.0 | 24.9 | 27.0 |
| Chronic conditions (yes)b | % | 28.8 | 32.5 | 35.2 | 34.1 | 31.6 |
| Depressive symptoms (yes)c | % | 7.6 | 7.3 | 3.7 | 7.5 | 2.8 |
| Total energy intake (kcal, day) | mean (se) | 2115.6 (10.5) | 2269.6 (73.0) | 2223.2 (58.4) | 2154.5 (46.6) | 2294.1 (66.0) |
| Total sugar intake (gram, day) | mean (se) | 113.2 (0.9) | 124.0 (4.9) | 134.1 (6.3) | 122.3 (3.5) | 128.7 (55) |
| Total chocolate intake (gram, day) | mean (se) | n.a | 18.7 (0.5) | 60.5 (1.1) | 100 (0.0) | 196.9 (5.6) |
| Total dark chocolate intake (gram, day) | mean (se) | n.a | 0.01 (0.04) | 0.34 (0.03) | 0.06 (0.02) | 0.09 (0.02) |
| a. All estimated are weighted to be nationally representative |
| b. Depression symptoms measured using Patient Health Questionnaire (PHQ-9): None or minimum to mild (0-9) Moderate to severe (10-27) |
| c. Chronic conditions include cardiovascular disease, arthritis, and cancer. |